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Boneberg, Franziska M ; Brandmann, Tobias ; Kobel, Lena ; van den Heuvel, Jasmin ; Bargsten, Katja  
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# Molecular mechanism of the RNA helicase DHX37 and its activation by UTP14A in ribosome biogenesis

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## ABSTRACT

Eukaryotic ribosome biogenesis is a highly orchestrated process involving numerous assembly factors including ATP-dependent RNA helicases. The DEAH helicase DHX37 (Dhr1 in yeast) is activated by the ribosome biogenesis factor UTP14A to facilitate maturation of the small ribosomal subunit. We report the crystal structure of DHX37 in complex with single-stranded RNA, revealing a canonical DEAH ATPase/helicase architecture complemented by a structurally unique carboxy-terminal domain (CTD). Structural comparisons of the nucleotide-free DHX37-RNA complex with DEAH helicases bound to RNA and ATP analogs reveal conformational changes resulting in a register shift in the bound RNA, suggesting a mechanism for ATP-dependent 3'–5' RNA translocation. We further show that a conserved sequence motif in UTP14A interacts with and activates DHX37 by stimulating its ATPase activity and enhancing RNA binding. In turn, the CTD of DHX37 is required, but not sufficient, for interaction with UTP14A *in vitro* and is essential for ribosome biogenesis *in vivo*. Together, these results shed light on the mechanism of DHX37 and the function of UTP14A in controlling its recruitment and activity during ribosome biogenesis.

**Keywords:** RNA helicase; X-ray crystallography; ribosome biogenesis

## INTRODUCTION

Composed of four ribosomal RNAs (rRNAs) and ~80 proteins, eukaryotic ribosomes are complex ribonucleoprotein (RNP) nanomachines whose biogenesis is a highly orchestrated and regulated process (Kressler et al. 2017; Peña et al. 2017). Ribosome assembly starts in the nucleolus with the transcription of a large precursor, known as the 47S preribosomal RNA (pre-rRNA) in mammalian cells, containing the future 18S, 5.8S and 28S rRNAs interspaced internally with noncoding internal transcribed spacers (ITS1 and ITS2) and flanked by 5' and 3' external transcribed spacers (5' and 3' ETS). The pre-rRNA transcript undergoes extensive processing and modification concurrently with RNA folding and ribosomal protein recruitment (Henras et al. 2015). An early cleavage event within ITS1 separates the 18S rRNA and 5.8S-28S rRNA fragments, leading to independent assembly pathways of the 40S and 60S subunits, which are eventually exported to the cytoplasm for their final maturation. Eukaryotic ribosome biogenesis is dependent on ~75 small nucleolar RNAs (snoRNAs) that guide rRNA folding and site-specific mod-

ifications, and more than 200 protein assembly factors, including exo- and endonucleases, ATPases, GTPases, and DExH/D-box RNA helicases (Woolford and Baserga 2013). U3 small nucleolar RNA (snoRNA) is a key component of the small subunit (SSU) processome, an early stable intermediate of 40S subunit assembly (Phipps et al. 2010). By hybridizing to the 5'-ETS and regions of the pre-18S rRNA, U3 facilitates initial folding of the pre-rRNA and formation of individual 18S rRNA domains, and is required for nucleolytic cleavage within the 5' ETS and ITS1 (Beltrame and Tollervey 1992).

DExH/D-box ATPases belonging to the SF2 helicase superfamily use their ATP-dependent RNA binding or translocation activities to catalyze structural remodeling of RNAs or RNP complexes (Jankowsky and Fairman 2007; Fairman-Williams et al. 2010; Jankowsky 2011; Jarmoskaite and Russell 2014). Whereas DEAD-box helicases unwind RNA secondary structures by binding to and distorting RNA duplexes in an ATP-dependent manner (Yang et al. 2007; Hilbert et al. 2009), DEAH/RHA helicases (also termed DExH or DEAH helicases, and referred to as DEAH helicases herein) generally unwind RNA duplexes

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or remodel RNPs by processive, ATP-driven translocation along an RNA strand in the 3′–5′ direction (Büttner et al. 2007; Pyle 2008; He et al. 2010; Walbott et al. 2010; Linder and Jankowsky 2011). DExH/D-box helicases comprise a core module built of two RecA-like domains that utilize conserved sequence and structural motifs to bind ATP (motifs I, II, III, Va, and VI) and RNA (motifs Ia, Ib, Ic, IV, IVa, and V) (Jankowsky and Fairman 2007). Most DExH/D-box helicases do not possess intrinsic substrate specificity since RNA binding occurs via the ribose-phosphate backbone (Sengoku et al. 2006; Jankowsky 2011; Ozgur et al. 2015). Instead, they rely on protein cofactors for recruitment to specific RNAs and catalytic activation (Sloan and Bohnsack 2018). The specificity and catalytic activity of DEAH helicases is typically regulated by interactions with G-patch proteins (Robert-Paganin et al. 2015).

In yeast, 17 DExH/D-box RNA helicases are essential for ribosome biogenesis as they promote snoRNA removal and effect structural rearrangements in the assembling ribosomal subunits (Martin et al. 2014). The DEAH helicase Dhr1 is required for the assembly of the 40S subunit (Colley et al. 2000; Granneman et al. 2006). Using its helicase activity, Dhr1 displaces U3 snoRNA from the 5′-terminal part of the pre-18S rRNA, thereby promoting the formation of the central pseudoknot, a critical step in the folding of the 18S rRNA (Sardana et al. 2015). Dhr1 is supported by the ribosomal biogenesis factor Utp14 and the rRNA methyltransferase-coactivator complex Bud23–Trm112, both of which physically interact with the helicase and facilitate its efficient association with the preribosomal particle (Sardana et al. 2014; Zhu et al. 2016). Previous studies have suggested that Utp14, a stable component of the SSU processome, activates the RNA unwinding activity of Dhr1 in vitro and is required for the association of Dhr1 with the preribosome in vivo (Zhu et al. 2016). This suggests that Utp14 regulates the recruitment and activity of Dhr1 during the transition from the SSU processome to the pre-40S particle, and thereby the timing of the maturation of the 40S subunit. Dhr1 and Utp14 have highly conserved mammalian orthologs, DHX37 and UTP14A, both of which have also been implicated in ribosome biogenesis (Choudhury et al. 2018). However, the structural and mechanistic details of the RNA remodeling activity of Dhr1/DHX37 and its stimulation by Utp14/UTP14A remain elusive at present.

To obtain insights into the molecular mechanism of DHX37/Dhr1 helicases in ribosome biogenesis, we determined the crystal structure of murine DHX37 in complex with single-stranded RNA in the absence of adenosine nucleotide, which reveals a conformational state likely corresponding to a nucleotide exchange intermediate and suggests a mechanism for ATP-dependent 3′–5′ RNA translocation. We further demonstrate that UTP14A stimulates both the ATPase and RNA binding activities of DHX37, thereby enhancing the processivity of the helicase.

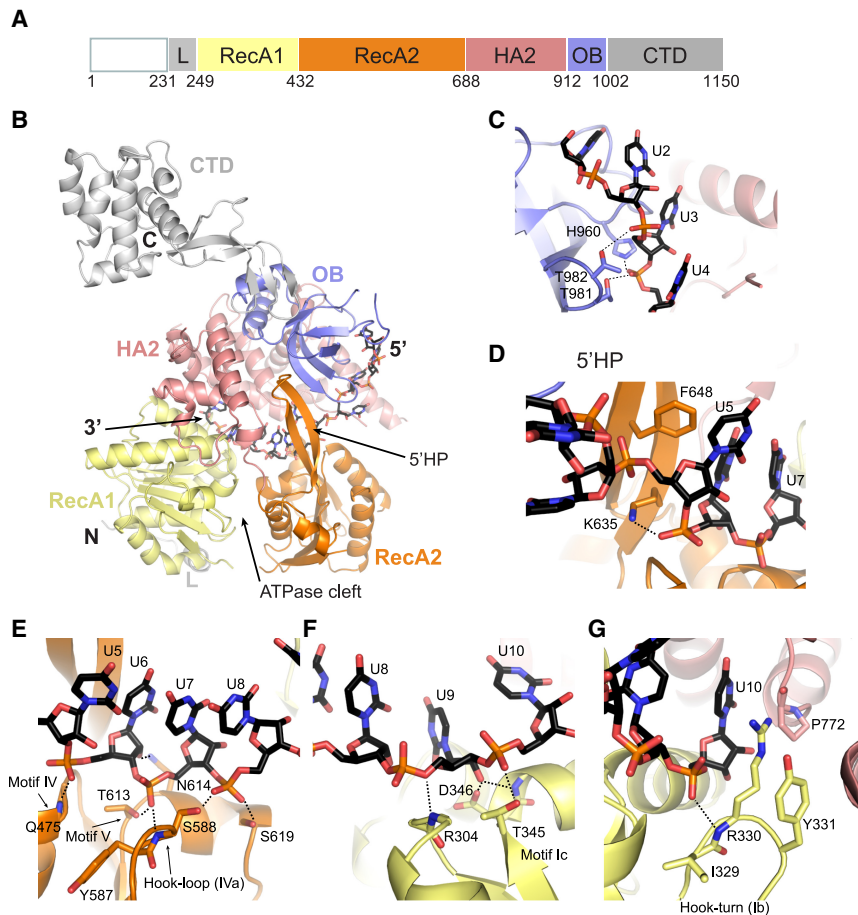
Finally, we show that a structurally unique carboxy-terminal domain (CTD) of DHX37, absent from other DEAH helicases, is required for the interaction with UTP14A in vitro and the function of DHX37 in vivo. Combined, these insights shed light on the role of UTP14A in controlling the recruitment and activation of DHX37 during 40S subunit maturation.

## RESULTS AND DISCUSSION

### Structure of DHX37 bound to ssRNA in the absence of adenosine nucleotide

DHX37 is a ~130 kDa multidomain protein containing a canonical DEAH ATPase/helicase catalytic module that comprises two RecA-like (RecA1 and RecA2) domains and a helicase-associated domain (HA2), followed by an oligonucleotide/oligosaccharide-binding fold (OB) domain (Fig. 1A; Ozgur et al. 2015). The RecA1 domain is preceded by an extended region predicted to lack tertiary structure. Aiming to obtain structural insights into the molecular mechanism of DHX37, we expressed an amino-terminally truncated variant of murine DHX37 (MmDHX37) comprising amino acid residues 179–1150 (MmDHX37<sup>179–1150</sup>). The RNA-dependent ATPase activity of MmDHX37<sup>179–1150</sup> was indistinguishable from that of full-length MmDHX37, indicating that the amino-terminal region of DHX37 is not required for catalytic activity (Supplemental Fig. S1A; Supplemental Table S1). Fluorescence-polarization assays were used to monitor RNA binding, revealing that MmDHX37<sup>179–1150</sup> exhibited highest affinity for oligouridine RNA (Supplemental Fig. S1B). RNA binding was reduced in the presence of ADP or the nonhydrolyzable ATP analog AMPPNP (Supplemental Fig. S1C), indicating that DHX37 does not require adenosine nucleotides (ADP or ATP) for ssRNA binding and suggesting that RNA binding is weakened in the ATP- or ADP-bound states, as observed for other processive SF2 helicases (Levin et al. 2003; Robert-Paganin et al. 2016). This was corroborated by a pull-down experiment utilizing biotin-coupled oligouridine RNA (Supplemental Fig. S1D), which confirmed that stable ssRNA binding is independent of adenosine nucleotides.

MmDHX37<sup>179–1150</sup> was subsequently crystallized in complex with uridine decanucleotide (U<sub>10</sub>), and its structure determined by single-wavelength anomalous X-ray diffraction experiment. The structure was refined at a resolution of 2.9 Å to an  $R_{\text{work}}$  of 26.8% and  $R_{\text{free}}$  of 29.7%. The structure reveals a canonical DEAH ATPase/helicase architecture (Fig. 1B), in which the RecA-like, HA2, and OB-fold domains form a compact core, as previously observed in the structures of the related DEAH/RHA helicases Prp43 (He et al. 2017; Tauchert et al. 2017), MLE (Prabu et al. 2015), DHX36 (Chen et al. 2018b,c), and Prp2 (Schmitt et al. 2018). An additional  $\alpha$ -helical carboxy-terminal



**FIGURE 1.** Structure of the DHX37-RNA complex in adenosine nucleotide-free state. (A) Schematic of the domain architecture of MmDHX37. The crystallized protein comprises residues 179–1150. (L) linker domain, (RecA1) RecA-like domain 1, (RecA2) RecA-like domain 2, (HA2) helicase-associated domain 2, (OB) Oligonucleotide-binding fold domain, (CTD) carboxy-terminal domain. (B) Overall structure of the MmDHX37<sup>179–1150</sup>-U<sub>10</sub> RNA complex. The helicase is shown in cartoon format and colored according to the scheme in A. The bound RNA is colored black and shown in stick format. The amino-/carboxy- and 5'/3'-termini of the DHX37 polypeptide and the RNA are indicated. 5'HP: 5' hairpin motif. (C) Zoom-in view of RNA nucleotides U1–U4. Hydrogen bonding interactions are indicated with dashed black lines. (D) Zoom-in view of the 5'HP and RNA nucleotides U3–U7. (E) Zoom-in view of nucleotides U5–U8 binding to helicase motifs IV, Iva, and V. (F) Zoom-in view of nucleotides U8–U10 binding to helicase motif Ic. (G) Zoom-in view of nucleotides U9–U10 binding to helicase motif Ib (hook-turn).

domain (CTD, Fig. 1A), a distinct feature of DHX37 orthologs, projects away from the helicase core (Fig. 1B). The CTD has a unique fold not observed in other proteins and makes few contacts with the helicase core, its orientation being maintained by crystal contacts with a second DHX37 molecule in the asymmetric unit of the crystal. In solution, the CTD is thus likely to be flexibly tethered to the helicase core and conformationally dynamic.

Although the MmDHX37<sup>179–1150</sup>-U<sub>10</sub> complex was crystallized in the presence of the nonhydrolyzable ATP analog AMPPNP, no corresponding electron density could be observed in the ATPase active site located at the interface of the RecA1 and RecA2 domains. Structural comparison of

the DHX37-U<sub>10</sub> complex with that of *Chaetomium thermophilum* Prp43 bound to U<sub>7</sub> RNA and the transition state analog ADP.BeF<sub>3</sub> (Tauchert et al. 2017) reveals that the DHX37 ATPase active site has an open conformation in which conserved active site motifs involved in ATP and Mg<sup>2+</sup> coordination (Supplemental Fig. S3) adopt conformations incompatible with high-affinity ATP binding (Supplemental Fig. S4). In particular, superposition of the RecA1 domains of the DHX37 and Prp43 complexes shows that active site motifs Va and VI in the DHX37 RecA2 domain are displaced by ~7 Å away from the position that they assume in the ATP-bound conformation in Prp43 (Supplemental Fig. S4C) and other DEAH helicases (Prabu et al. 2015; He et al. 2017; Tauchert et al. 2017; Schmitt et al. 2018).

The U<sub>10</sub> RNA is bound predominantly by sequence nonspecific ionic and hydrogen-bonding interactions with the ribose-phosphate backbone (Supplemental Fig. S2B), as observed for other DEAH helicases (Prabu et al. 2015; He et al. 2017; Tauchert et al. 2017; Chen et al. 2018b,c). DHX37 makes base-directed interactions only with nucleotides U1, U5 and U9 (Supplemental Fig. S5), which may contribute to the observed binding preference of DHX37 for uridine-rich RNA (Supplemental Fig. S1B). The 5'-terminal part of the RNA binds in a surface cleft along the OB domain, with phosphate groups of nucleotides U3 and U4 contacted by two threonine residues in the β20–β21 loop in the OB domain and by His960 in the

β18–β19 loop (Fig. 1C). In turn, nucleotides U5–U10 are bound in a central channel within the helicase core, surrounded by the RecA1, RecA2, HA2, and OB domains. Throughout the channel, the RNA is stabilized by ionic interactions with the basic side chain groups of phylogenetically conserved amino acid residues Lys635 (RecA2 domain), Arg803 (HA2), Arg303, Arg304, and Arg330 (all RecA1). At the 5'-proximal end of the RNA-binding channel, the RNA backbone passes through a narrow opening formed by the HA2 and OB domains and a long β-hairpin that extends from the RecA2 domain and contacts the HA2 and OB domains. The β-hairpin (5' HP), which is a characteristic feature of the DEAH helicases (He et al. 2017), packs



against nucleotide U5 forming a 5'-terminal bookend for the RNA nucleotides bound in the helicase channel (Fig. 1D). The 5'HP may additionally provide a physical barrier to induce the unwinding of secondary structures in RNA or remodeling of protein-RNA interactions during 3'-5' translocation, as has been proposed for other SF2 superfamily RNA helicases (Büttner et al. 2007; Pyle 2008; Ozgur et al. 2015; He et al. 2017; Tauchert et al. 2017). Five nucleotides (U6-U10) are accommodated in the central part of the RNA binding channel. The bases of nucleotides U6-U10 form a continuous stack, while their ribose-phosphate backbone makes numerous contacts with the RNA binding motifs IV, V, and VI in the RecA2 domain and motifs Ib and Ic in the RecA1 domain (Fig. 1E-G). Specifically, motif IV contacts the phosphate group of U6 via hydrogen-bonding interaction with the backbone amide of Gln475. An invariant threonine residue in motif V (Thr613) interacts with the phosphate group of U7. Motif IVa, also described as the hook-loop (Prabu et al. 2015), binds the phosphate groups of U7 and U8 by hydrogen-bonding interactions with the backbone amide and side chain groups of Ser588, respectively (Fig. 1E). In the 3'-proximal part of the RNA-binding channel, the ribose-phosphate moieties of nucleotides U9 and U10 interact with the RecA1 domain. The backbone amide group of Arg304 makes a hydrogen-bonding interaction with the phosphate group of U9, while the 2'-hydroxyl group of U9 is contacted by the side-chain carboxyl group of Asp346 (Fig. 1F). The phosphate group of U10 is bound by hydrogen-bonding interactions with the invariant residue Thr345 of motif Ic and the backbone amide of Arg330 of motif Ib (Fig. 1F), also referred to as the hook-turn (Tauchert et al. 2017). Arg330 and Tyr331 in motif Ib, together with invariant residue Pro772 in the  $\alpha 15$ - $\alpha 16$  loop of the HA2 domain, line the 3'-terminal exit of the RNA binding channel, forming a 3'-terminal bookend for the bound RNA (Fig. 1G).

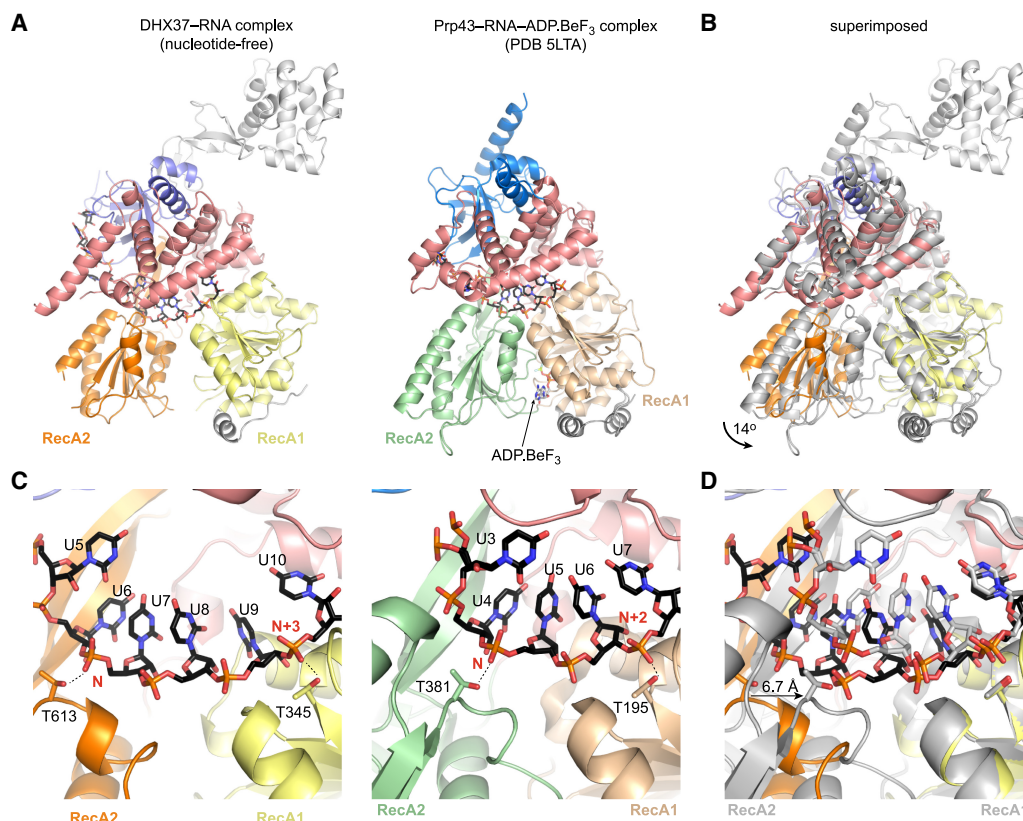
Altogether, the structure of the DHX37-U<sub>10</sub> complex reveals the RNA binding mode of DHX37 in the absence of adenosine nucleotide (ATP or ADP), which likely corresponds to a nucleotide exchange state of the helicase upon the dissociation of ATP hydrolysis products (ADP and inorganic phosphate) and prior to the binding of the next ATP molecule in the catalytic cycle. The conformation of the DHX37 helicase core and its RNA binding mode bears resemblance to the structures of the related helicase DHX36 bound to a DNA substrate in the absence of ATP (Chen et al. 2018b,c), suggesting that the ATP-free conformation is conserved across the DEAH helicase family (Supplemental Fig. S6).

### RNA binding mode reveals the 3'-5' translocation mechanism of DHX37

DEAH RNA helicases are generally thought to catalyze the unwinding of RNA secondary structures or remodeling of

protein-RNA complexes by ATP-powered translocation along a tracking RNA strand in the 3'-5' direction (Ozgur et al. 2015). In this process, the helicases cycle through ATP-, ADP-bound, and nucleotide-free conformational states, in which ATP nucleotide binding at the interface of the RecA1 and RecA2 domains allosterically induces structural transitions in the RNA binding sites. Although the translocation directionality of DHX37 (or Dhr1) has not been formally demonstrated, Prp43, a close homolog of DHX37 within the DEAH family (~39% sequence identity within the helicase cores of human DHX37 and DHX15/Prp43), is a bona fide 3'-5' helicase (He et al. 2017). Structural superpositions of MmDHX37<sup>179-1150</sup> in the nucleotide-free state with Prp43 in complexes with oligo-U RNAs and ATP analogs reveal major conformational differences in the relative orientations of the RecA1 and RecA2 domains, reflecting an open conformation of the ATPase site in DHX37 and a closed conformation in ATP analog-bound Prp43 (He et al. 2017; Tauchert et al. 2017). Structural analysis using the DynDom server (Hayward and Berendsen 1998) further indicates that the RecA2 domain of DHX37 must rotate by ~14° with respect to RecA1 in order to assume the ATP-bound conformation, while the HA2 and OB domains undergo a minor ~2° rotation combined with a 2 Å shift relative to RecA1 (Fig. 2A,B). The conformational differences between the ATP-free and ATP analog-bound states are reflected in distinct RNA binding modes observed in the complexes. Whereas DHX37 in the ATP-free state encloses a 5-nt stack within the RNA binding tunnel, Prp43 in the ATP-bound state accommodates a 4-nt stack (Fig. 2C,D). In both states, the 5' and 3' termini of the nucleotide stacks are supported by the 5'HP and motif Ib hook-turn, respectively. As a consequence of the open conformation of the RecA1 and RecA2 domains, the RNA-interacting motifs in the RecA2 domain, including motif V containing Thr613, shift by one nucleotide toward the 5' end of the RNA (Fig. 2D). Conversely, while Thr613 remains in contact with the phosphate moiety of nucleotide N, the invariant Thr345 of motif Ic that interacts with the phosphate of nucleotide N + 3 in the ATP-free state would contact nucleotide N + 2 in the ATP-bound state upon closure of the RecA1 and RecA2 domains (Fig. 2C).

Together, the structural insights provided here suggest a model for ATP-dependent 3'-5' translocation in DHX37 and other DEAH helicases, in which ATP binding at the interface of the RecA1 and RecA2 domains induces a conformational closure that is propagated to the RNA binding channel. By maintaining interactions with motifs IV, IVa and V and the 5'HP in RecA2, the RNA is shifted by one nucleotide along RecA1 and the 3'-terminal nucleotide exits past the hook-turn motif. Upon ATP hydrolysis and ADP/Pi release, the helicase reverts to the open conformation, which allows the RecA2 domain to undergo a register shift and engage the next 5'-terminal nucleotide, while motifs Ic and Ib (hook-turn) in RecA1 remain tightly bound to the



**FIGURE 2.** RNA binding mode of DHX37 suggests a mechanism for ATP-driven 3′–5′ translocation. (A) Side-by-side comparison of the structures of the DHX37–RNA (*left*) and Prp43–RNA–ADP.BeF<sub>3</sub><sup>−</sup> (PDB entry 5LTA, *right*) complexes. The helicases were superimposed using their RecA1 domains and are shown in identical orientations. (B) Superposition of DHX37 (colored as in A) and Prp43 (colored gray). The ADP.BeF<sub>3</sub><sup>−</sup> transition state mimic is shown in stick format. (C) Side-by-side views of the RNA binding channels in the DHX37–RNA (*left*) and Prp43–RNA–ADP.BeF<sub>3</sub><sup>−</sup> (*right*) complexes. While DHX37 accommodates a 5-nt stack (U6–U10) in the ATP-free state, Prp43 accommodates a 4-nt stack (U4–U7) in the ADP.BeF<sub>3</sub><sup>−</sup> bound state. Motif IVa is omitted from the figures to give unobstructed view of the bound RNA. (D) Superposition of the RNA binding channels in DHX37 (colored as in A) and Prp43 (gray).

3′-terminal part of the RNA. In this mechanism, the helicase toggles between states in which 4- and 5-nt RNA stacks are accommodated in the RNA binding channel, which would be consistent with an “expansion” model of helicase translocation proposed by earlier studies (Supplemental Fig. S7; Pyle 2008; He et al. 2017; Chen et al. 2018a; Hamann et al. 2019). Notably, the model posits that ATP binding, rather than hydrolysis, provides the power stroke for translocation. This might explain previous observations that under presteady state conditions, yeast Dhr1 is able to unwind short RNA duplexes in an ATP-dependent manner but in the absence of actual ATP hydrolysis, since translocation along a single nucleotide might be sufficient to induce duplex unwinding in this case (Sardana et al. 2015).

### A short motif in UTP14A activates DHX37 by stimulating its ATPase activity and enhancing RNA binding

In yeast, the ribosome biogenesis factor Utp14 associates with Dhr1, the ortholog of DHX37, recruiting it to the peri-

bosomal particle and stimulating its RNA helicase activity to facilitate the dissociation of U3 snoRNP from the maturing 40S ribosomal subunit (Zhu et al. 2016). Utp14 directly interacts with Dhr1 in vitro and enhances its RNA unwinding activity (Zhu et al. 2016). To investigate whether the mammalian ortholog UTP14A has a similar effect on DHX37, we coexpressed MmDHX37<sup>179–1150</sup> together with full-length human UTP14A in baculovirus-infected insect cells and succeeded in purifying the resulting complex, confirming a direct physical interaction between DHX37 and UTP14A (Supplemental Fig. S8). We subsequently measured the ATPase activity of isolated MmDHX37<sup>179–1150</sup> and the MmDHX37<sup>179–1150</sup>–UTP14A complex. At physiological ATP concentrations (2 mM), the ATPase activity of the MmDHX37<sup>179–1150</sup>–UTP14A complex was ~15-fold higher than that of MmDHX37<sup>179–1150</sup> alone (Supplemental Fig. 9A,B; Supplemental Table S1), suggesting that UTP14A functions as an activator of DHX37 in part by increasing its RNA-dependent ATPase activity. This result stands in contrast to a previous observation that yeast Utp14 increases the RNA unwinding activity of Dhr1 in vitro

without affecting ATPase activity (Zhu et al. 2016) but is in good agreement with a recent study of human DHX37 (Choudhury et al. 2018). Although DHX37 and Dhr1 have similar biological functions, it is possible that they use divergent molecular mechanisms.

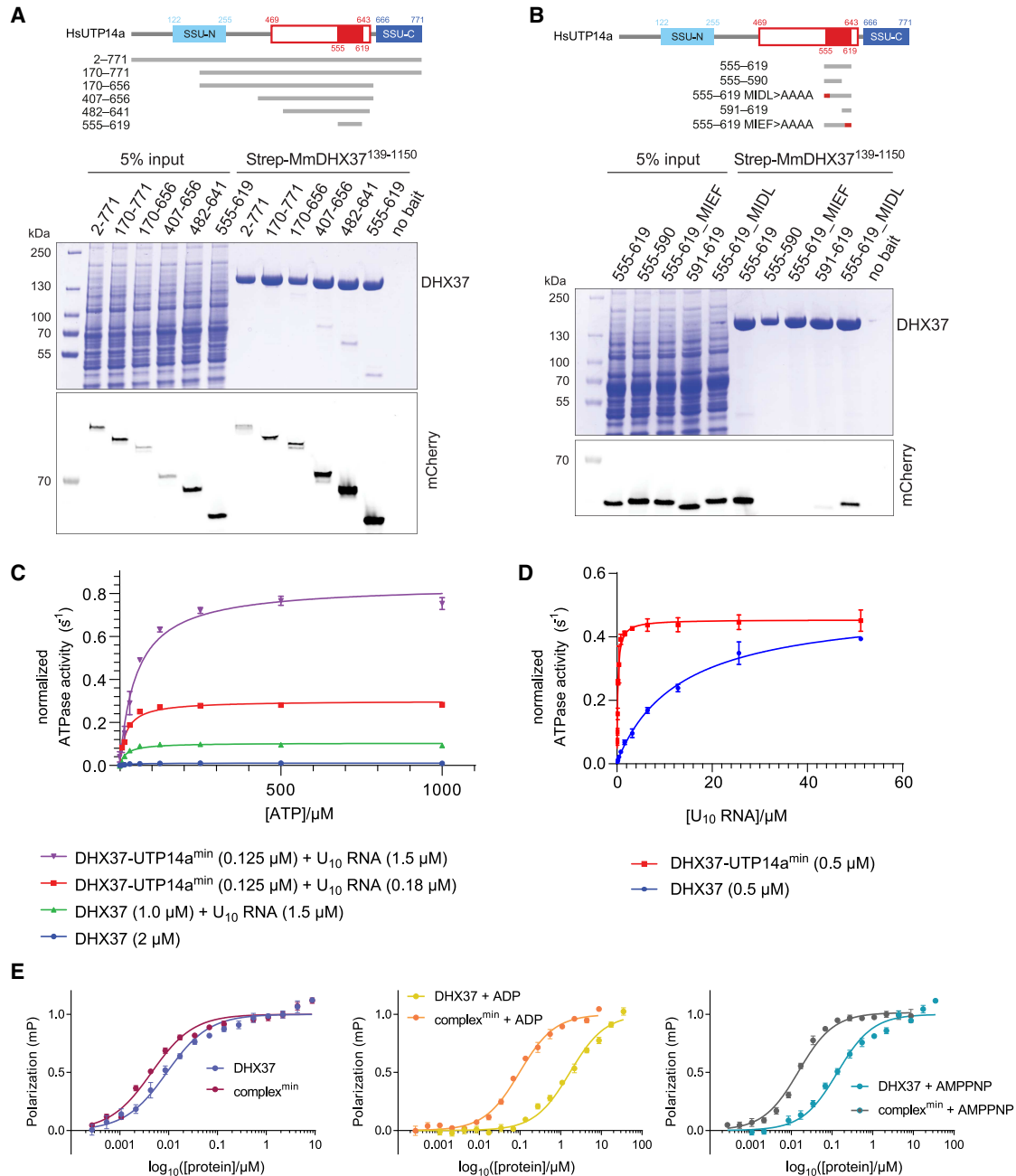
Recent structural studies of the yeast SSU processome showed that Utp14 is an intrinsically unfolded, extended protein whose amino-terminal and carboxy-terminal regions participate in SSU processome assembly by making extensive contacts with other ribosome biogenesis factors including Utp6, Sof1, and Utp7 (Barandun et al. 2017; Cheng et al. 2017; Black et al. 2018). The intervening region (Supplemental Fig. S10A), which is disordered in the SSU processome structures, has been shown to be necessary for interaction with Dhr1 (Zhu et al. 2016). To define the region in human UTP14A responsible for interaction with DHX37, we expressed a series of truncated protein constructs in human cells and tested their binding to immobilized MmDHX37<sup>139–1150</sup> in a pull-down assay. A UTP14A protein fragment comprising residues Lys555–Ser619 was efficiently coprecipitated by MmDHX37<sup>139–1150</sup> (Fig. 3A), while further amino- and carboxy-terminal truncations abolished binding (Fig. 3B), suggesting that this region (hereafter referred to as UTP14A<sup>min</sup>) is the minimal motif necessary and sufficient for interaction with DHX37. Purified recombinant UTP14A<sup>min</sup> protein was also sufficient to activate DHX37 in vitro, enhancing its ATPase activity by a factor of ~14, indicating that the stimulatory effect can be attributed solely to the minimal region necessary for physical interaction with DHX37 (Supplemental Fig. S9C,D; Supplemental Table S1). UTP14A<sup>min</sup> contains two conserved sequence motifs; the amino-terminal motif (Met558–Leu561) is conserved in metazoan DHX37 orthologs, while the carboxy-terminal motif (Met594–Glu597) is highly conserved across all eukaryotes including yeast (Supplemental Fig. S10B). Mutation of the amino-terminal motif residues to alanine reduced binding to MmDHX37<sup>139–1150</sup>, while mutation of the carboxy-terminal motif completely abolished interaction with MmDHX37<sup>139–1150</sup> (Fig. 3B). Accordingly, the carboxy-terminal motif mutant UTP14A<sup>min</sup> protein failed to stimulate the RNA-dependent ATPase activity of MmDHX37<sup>179–1150</sup> in vitro, while the activating effect of the amino-terminal motif mutant protein was substantially reduced (Supplemental Fig. 9C,D). Together, these experiments demonstrate that a conserved motif in the central region of UTP14A physically interacts with DHX37 and allosterically activates its ATPase activity. This is consistent with prior genetic experiments showing that point mutations and deletions in the corresponding region of yeast Utp14 perturb its interaction with Dhr1 and impair the function of Dhr1 in ribosome biogenesis (Zhu et al. 2016). Although UTP14A orthologs contain a highly conserved glycine-rich sequence (Gly630–Gly638 in human UTP14A, Supplemental Fig. S10B) somewhat reminiscent

of the G-patch protein consensus motif, our deletion analysis indicates that this sequence in UTP14A is neither required for the interaction with DHX37 nor for the enhancement of its ATPase activity. This suggests that UTP14A is not a canonical G-patch protein, as proposed previously (Zhu et al. 2016).

To investigate the mechanism underpinning UTP14A-mediated activation of DHX37, we analyzed the ATPase activities of MmDHX37<sup>179–1150</sup> and the MmDHX37<sup>179–1150</sup>–UTP14A<sup>min</sup> complex as a function of ATP and RNA concentrations. UTP14A<sup>min</sup> enhanced the ATPase activity of DHX37 in the presence of 1.5  $\mu$ M U<sub>10</sub> RNA by increasing  $k_{cat}$  by approximately eightfold, while simultaneously reducing the Michaelis constant ( $K_M$ ) for ATP by ~2.7-fold (Table 1; Fig. 3C). The half-maximal effective concentration ( $EC_{50}$ ) of U<sub>10</sub> RNA was reduced by a factor of ~60-fold in the presence of UTP14A<sup>min</sup> (Table 1; Fig. 3D). Together, these results suggest that UTP14A-mediated stimulation of the RNA-dependent ATPase activity of DHX37 is in large part due to enhancement of RNA binding. To corroborate our kinetics measurements, we examined the ssRNA binding activities of MmDHX37<sup>179–1150</sup> and the MmDHX37<sup>179–1150</sup>–UTP14A<sup>min</sup> complex in the absence and presence of adenosine nucleotides using fluorescence polarization (FP) (Fig. 3E). In the absence of adenosine nucleotides, UTP14A<sup>min</sup> stimulated the binding of U<sub>10</sub> RNA by approximately twofold. However, in the presence of ADP or AMPPNP, the MmDHX37<sup>179–1150</sup>–UTP14A<sup>min</sup> complex displayed a ~25-fold or ~10-fold higher affinity, respectively, for U<sub>10</sub> than MmDHX37<sup>179–1150</sup> alone (Fig. 3E). This indicates that the interaction with UTP14A increases the nucleotide-dependent RNA binding activity of DHX37 and that the increase is commensurate with the observed enhancement of ATPase activity engendered by UTP14A (Supplemental Fig. S9). Collectively, these results imply that UTP14A stimulates the activity of DHX37 by functioning as a processivity factor that enhances nucleotide-dependent RNA binding, thereby increasing the mechanochemical coupling between ATP binding/hydrolysis and RNA translocation. This mechanism thus likely underpins the regulation of the ATP-dependent RNA translocation activity of DHX37, which is essential for the remodeling of the U3 snoRNA-pre-18S rRNA structure during 40S maturation.

### The carboxy-terminal domain of DHX37 is essential for UTP14A binding in vitro and ribosome biogenesis in vivo

G-patch proteins activate DEAH RNA helicases by interactions with the OB domain (Tanaka and Schwer 2005; Lebaron et al. 2009; Chen et al. 2014; Robert-Paganin et al. 2015; Heininger et al. 2016). As prior studies (Zhu et al. 2016) and our biochemical experiments suggested that UTP14A is functionally distinct from canonical G-patch



**FIGURE 3.** UTP14A stimulates ATPase activity of DHX37 and enhances RNA binding. (A, top) Schematic diagram of human UTP14A and constructs used for pull-down experiment. SSU-N and SSU-C denote the amino- and carboxy-terminal regions corresponding to the regions of Utp14 bound in the yeast SSU processome structure. Red boundary denotes region corresponding to the Dhr1 interacting region in yeast Utp14. Solid red box denotes minimal region in HsUTP14A required for interaction with DHX37. (Bottom) SDS-PAGE analysis of streptactin pull-down experiment using StreptII-tagged MmDHX37<sup>139-1150</sup> as bait and indicated mCherry-tagged HsUTP14A constructs as prey. UTP14A constructs are visualized by mCherry fluorescence. (B, top) Schematic diagram of human UTP14A and constructs used for pull-down experiment. (Bottom) SDS-PAGE analysis of streptactin pull-down experiment using StreptII-tagged MmDHX37<sup>139-1150</sup> as bait and indicated mCherry-tagged HsUTP14A constructs as prey. MIEF denotes the M594A/I595A/E597A/F599A quadruple point mutant; MIDL denotes the M558A/I559A/D560A/L561A mutant. (C) RNA-dependent ATPase activity of MmDHX37<sup>179-1150</sup> or the MmDHX37<sup>179-1150</sup>-HsUTP14A<sup>min</sup> complex as a function of ATP concentration. The data were subjected to Michaelis-Menten analysis and the calculated  $k_{cat}$  and  $K_M$  values are provided in Table 1. Experiments were performed in triplicates, with data points indicating the mean and error bars indicating the standard error of the mean. (D) RNA-dependent ATPase activity of MmDHX37<sup>179-1150</sup> or the MmDHX37<sup>179-1150</sup>-HsUTP14A<sup>min</sup> complex as a function of U<sub>10</sub> RNA concentration. Experiments were performed in triplicates, with data points indicating the mean and error bars indicating the standard error of the mean. EC<sub>50</sub> values are provided in Table 1. (E) Fluorescence-polarization equilibrium binding assays of MmDHX37<sup>179-1150</sup> or the MmDHX37<sup>179-1150</sup>-HsUTP14A<sup>min</sup> complex with ATTO532 fluorophore-labeled U<sub>12</sub> RNA in the absence of adenosine nucleotides (left) and in the presence of ADP (center) and the nonhydrolyzable ATP analog AMPPNP (right).  $K_d$  values are provided in Table 1.



**TABLE 1.** Kinetic and thermodynamic parameters of DHX37 and the DHX37-UTP14A complex

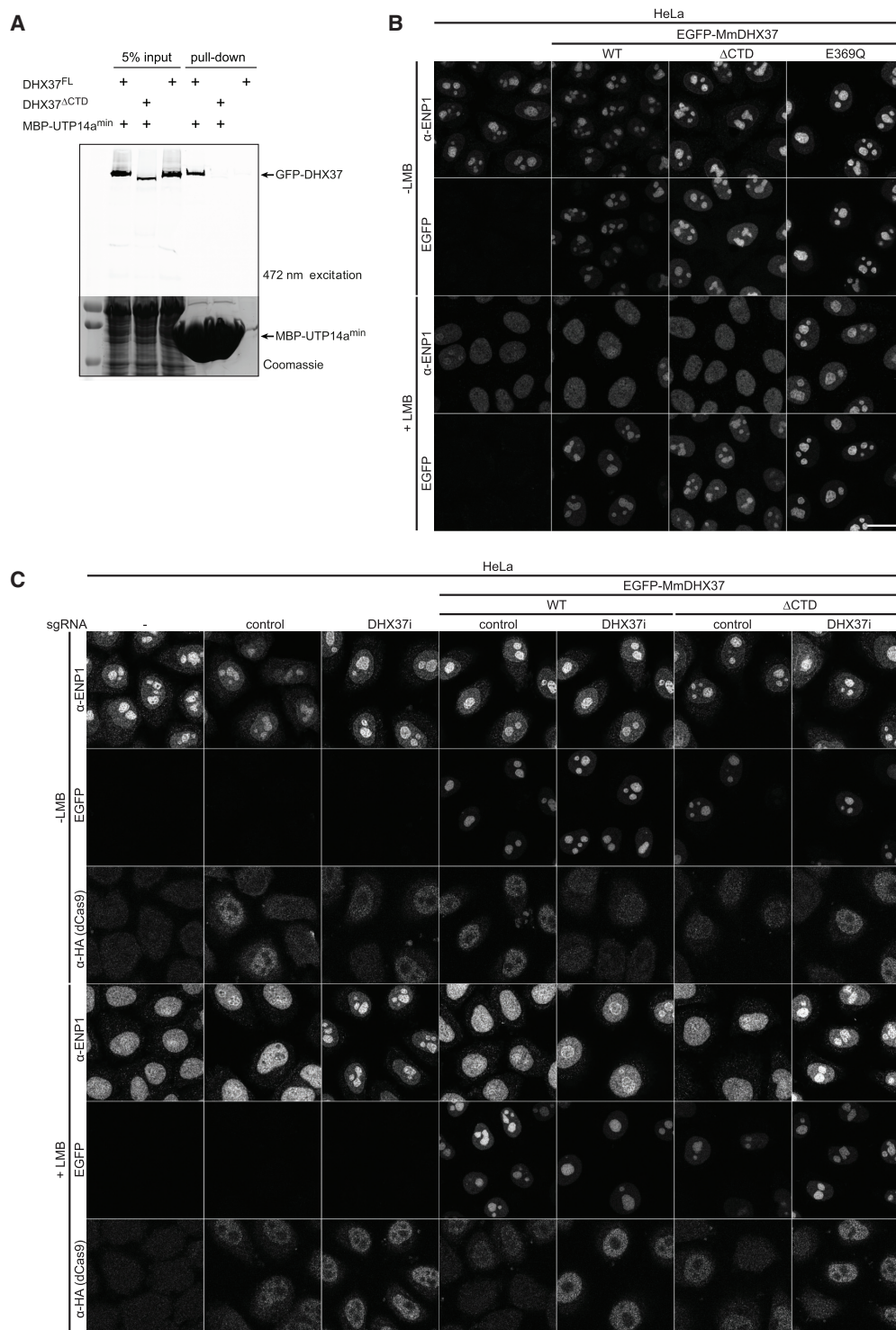
Enzyme	RNA	$K_m$ ( $\mu$ M)	$k_{cat}$ ( $s^{-1}$ )
ATPase activity			
MmDHX37 <sup>179–1150</sup>	—	28.30 $\pm$ 2.92	0.0108 $\pm$ 0.0001
MmDHX37 <sup>179–1150</sup> (1.0 $\mu$ M)	U <sub>10</sub> (1.5 $\mu$ M)	19.26 $\pm$ 4.79	0.105 $\pm$ 0.003
MmDHX37 <sup>179–1150</sup> + UTP14A <sup>min</sup> (0.125 $\mu$ M)	U <sub>10</sub> (1.5 $\mu$ M)	52.61 $\pm$ 4.78	0.844 $\pm$ 0.009
MmDHX37 <sup>179–1150</sup> + UTP14A <sup>min</sup> (0.125 $\mu$ M)	U <sub>10</sub> (0.188 $\mu$ M)	19.95 $\pm$ 0.75	0.301 $\pm$ 0.001
Enzyme	EC <sub>50</sub> ( $\mu$ M)		
EC <sub>50</sub> of U <sub>10</sub> RNA for ATPase activation (ATPase activity measured at 1 mM ATP)			
MmDHX37 <sup>179–1150</sup>	13.27 $\pm$ 1.08		
MmDHX37 <sup>179–1150</sup> + UTP14A <sup>min</sup>	0.22 $\pm$ 0.03		
Enzyme	Nucleotide	$K_d$ (nM)	
Equilibrium dissociation constants for U <sub>10</sub> RNA binding			
MmDHX37 <sup>179–1150</sup>	—	9.83 $\pm$ 0.55	
MmDHX37 <sup>179–1150</sup> + UTP14A <sup>min</sup>	—	4.19 $\pm$ 0.27	
MmDHX37 <sup>179–1150</sup>	ADP (1 mM)	2133 $\pm$ 316	
MmDHX37 <sup>179–1150</sup> + UTP14A <sup>min</sup>	ADP (1 mM)	86.06 $\pm$ 13.8	
MmDHX37 <sup>179–1150</sup>	AMPPNP (1 mM)	135.13 $\pm$ 2.41	
MmDHX37 <sup>179–1150</sup> + UTP14A <sup>min</sup>	AMPPNP (1 mM)	14.61 $\pm$ 0.26	

proteins, we next sought to pinpoint the specific domains in DHX37 involved in the interaction with UTP14A, focusing on the unique CTD of DHX37. We expressed full-length MmDHX37 and a truncated variant comprising amino acid residues 2–1003 (MmDHX37<sup>ΔCTD</sup>) in human cells and tested their binding to immobilized UTP14A<sup>min</sup> in a pull-down experiment. Whereas MmDHX37 was efficiently coprecipitated by UTP14A from cell lysate, MmDHX37<sup>ΔCTD</sup> was not, indicating that the CTD is required for stable interaction with UTP14A (Fig. 4A). The RNA-dependent ATPase activity of MmDHX37<sup>ΔCTD</sup> was similar to that of MmDHX37<sup>179–1150</sup> (Supplemental Fig. S11A). Crucially, the ATPase activity was not enhanced in the presence of UTP14A, indicating that the CTD is required both for the interaction with UTP14A and for UTP14A-mediated stimulation of the ATPase activity of DHX37 (Supplemental Fig. S11A). However, subsequent pull-down experiments revealed that purified recombinant DHX37 CTD (comprising amino acid residues 1003–1150) did not interact with UTP14A<sup>min</sup> (Supplemental Fig. S11B, C), indicating that although essential, the CTD is not sufficient to mediate a stable interaction with UTP14A.

To further test the requirement of the CTD for the function of DHX37 in ribosome biogenesis, we used a previously established reporter assay to examine the effect of overexpressing wild-type and mutant MmDHX37 proteins on the biogenesis of the 40S subunit. In this assay, nucleolar retention of the biogenesis factor ENP1 upon treatment with the nuclear export inhibitor LMB is indicative of an early 40S biogenesis defect (Badertscher et al. 2015). Overex-

pression of the ATPase-deficient E369Q point mutant resulted in nucleolar retention of ENP1, indicating that catalytically inactive DHX37 acts as a dominant-negative inhibitor of 40S biogenesis (Fig. 4B; Supplemental Fig. S11D). In contrast, overexpression of neither full-length wild-type MmDHX37 nor a catalytically active MmDHX37 lacking the CTD (MmDHX37<sup>ΔCTD</sup>) had a dominant-negative effect on 40S biogenesis. Notably, nucleolar localization of MmDHX37<sup>ΔCTD</sup> was not perturbed, indicating that the protein is folded and the CTD is not required for the nuclear import of DHX37 (Fig. 4B). To complement these results, we tested the ability of ectopically expressed full-length and MmDHX37<sup>ΔCTD</sup> proteins to rescue the 40S biogenesis defect caused by depletion of endogenous DHX37 in HeLa cells. To this end, we utilized CRISPR interference to target the promoter of the endogenous DHX37 gene, thereby down-regulating its expression, in cells stably transfected with MmDHX37 expression plasmids (Supplemental Fig. S11E). CRISPRi-targeted cells expressing MmDHX37<sup>ΔCTD</sup> exhibited nucleolar localization of ENP1 in the presence of LMB, indicating that MmDHX37<sup>ΔCTD</sup> was unable to rescue the early 40S biogenesis defect phenotype caused by DHX37 depletion, whereas expression of full-length MmDHX37 restored normal localization of ENP1 (Fig. 4C). Together, these observations indicate that the CTD of DHX37 is required for 40S biogenesis in vivo.

The current model for the function of DHX37/UTP14A, based on genetic and biochemical studies of yeast Dhr1/Utp14 (Zhu et al. 2016), posits that DHX37 is



**FIGURE 4.** The CTD of DHX37 is required for interaction with UTP14A in vitro and for ribosome biogenesis in vivo. (A) Pull-down experiment using recombinant maltose-binding protein-tagged UTP14A<sup>min</sup> and full-length and  $\Delta$ CTD MmDHX37 expressed in HEK293 cells. (B) Immunofluorescence analysis of the localization of the 40S subunit assembly factor ENP1 in HeLa cells expressing wild-type EGFP-MmDHX37, a truncated version lacking the carboxy-terminal domain ( $\Delta$ CTD), or the ATPase-deficient E369Q mutant. Expression of DHX37 proteins was induced with tetracycline for 24 h. Ninety minutes before fixation, cells were either supplemented with a solvent control or with 20 nM leptomycin B (LMB) to prevent CRM1-dependent nuclear export, leading to nucleoplasmic accumulation of ENP1 along with newly made 40S subunits in control cells (Badertscher et al. 2015). Note that the E369Q mutant prevents relocalization of ENP1 to the nucleoplasm in the presence of LMB, indicative of early 40S subunit maturation defects. (C) The CTD of DHX37 is required to rescue early 40S biogenesis defects upon DHX37 depletion. DHX37 was depleted from HeLa cells by CRISPRi. HeLa cells were left untreated (–) or transfected for 72 h with a plasmid encoding HA-tagged dCas9 and either a sgRNA targeting endogenous DHX37 (DHX37i) or a control sgRNA and transfected cells were selected with puromycin. Expression of the indicated EGFP-MmDHX37 rescue constructs was induced for 24 h with tetracycline. Cells were treated with LMB for 90 min before fixation as indicated. The expression of HA-tagged dCas9 and localization of the 40S assembly factor ENP1 were analyzed by immunofluorescence. Scale bars, 20  $\mu$ M.

recruited to the maturing 40S subunit in part by interactions with UTP14A and its ATP-dependent helicase activity is subsequently activated by UTP14A to promote the removal of U3 snoRNA, which triggers pre-18S rRNA folding and is required for ITS1 cleavage. In light of the results of the overexpression and rescue experiments, we conclude that the CTD of DHX37 is critical for this process. As with Dhr1 in yeast (Granneman et al. 2006; Sardana et al. 2015), overexpression of ATPase-deficient DHX37 inhibits 40S biogenesis, presumably because the helicase can still be recruited to the preribosomal particle but is unable to catalyze the removal of U3 snoRNA. Since DHX37 lacking the CTD is unable to exert the dominant-negative effect on 40S biogenesis, yet fails to rescue the phenotype caused by DHX37 depletion, our interpretation is that the CTD is required for UTP14A-dependent recruitment and activation of the helicase.

## Conclusions

RNA helicases, including DHX37/Dhr1, play indispensable roles in ribosome biogenesis. In this study, we determined the crystal structure of the DHX37-RNA complex in the nucleotide-exchange state, providing insights into the molecular mechanism for the ATP-dependent RNA translocation activity of DHX37 required for U3 snoRNA removal during 40S subunit maturation. Comparisons with structures of DHX36 and Prp43 in their respective nucleotide-free and the ATP-bound states suggest that ATP binding induces DHX37 to convert from an open conformation, in which a 5-nt RNA segment is accommodated in the RNA binding channel, to a closed conformation in which a 4-nt RNA segment is bound (Supplemental Fig. S7). Our structural insights thus lend further support to a mechanistic model for DEAH helicases in which conformational cycling between ATP-bound and nucleotide-free states results in directional 3' to 5' translocation along single-stranded RNA, which is coupled to RNA duplex unwinding or RNP remodeling.

Recent studies have suggested that UTP14A functions as a ribosome biogenesis factor in part by recruiting DHX37 to the maturing preribosomal particle (Choudhury et al. 2018). Our biochemical experiments reveal that a conserved motif in UTP14A, distinct from the canonical DEAH-interacting motif found in G-patch proteins, is necessary and sufficient for interaction with DHX37. The motif is part of the central region of UTP14A that is likely to be exposed on the surface of the SSU processome, poised to mediate DHX37 recruitment. In turn, interaction with UTP14A is mediated in part by the structurally unique CTD in DHX37, which is also required for ribosome biogenesis. Finally, our work suggests that UTP14A stimulates the activity of DHX37 by acting as a processivity factor that enhances RNA binding and the coupling between ATPase and RNA translocase/remodeling activities. Together,

these insights provide a mechanistic rationale for the coordination of DHX37 recruitment and activation in driving a key step in 40S subunit maturation and have implications for other DEAH helicases involved in ribosome biogenesis, such as Prp43. Determining the structural basis of the UTP14A–DHX37 interaction will be an important future step toward understanding the regulation of not only DHX37 but DEAH helicases in general.

## MATERIALS AND METHODS

### Protein expression and purification

All constructs used were cloned into MacroLab vectors developed by Scott Gradia (University of California, Berkeley) using ligation-independent cloning. Full-length murine DHX37 (MmDHX37) and an amino-terminally truncated fragment of murine DHX37 encompassing residues 179–1150 (MmDHX37<sup>179–1150</sup>) were inserted into the MacroLab vector 438B (Addgene #55219), in frame with an amino-terminal His<sub>6</sub>-tag cleavable using Tobacco Etch Virus (TEV) protease. Full-length human UTP14A (HsUTP14A) was inserted into the 438Rgfp vector (Addgene #55221) in frame with an amino-terminal (StreptII)-GFP-tag. The 438Rgfp-HsUTP14A construct was combined with 438B-MmDHX37<sup>179–1150</sup> into a single plasmid for subsequent coexpression in insect cells and copurification (Gradia et al. 2017). An amino-terminally truncated fragment of murine DHX37 encompassing residues 139–1150 (MmDHX37<sup>139–1150</sup>) was inserted into a modified 5B vector (Addgene #30122) with an amino-terminal His<sub>6</sub>-(StreptII)<sub>2</sub>-tag. The CTD of murine DHX37, encompassing residues 1003–1150 (MmDHX37<sup>CTD</sup>), was inserted into MacroLab vector 2G-T (Addgene #29707) in fusion with an amino-terminal His<sub>6</sub>-GST-tag. A truncated construct of human HsUTP14A encompassing residues 555–619 (HsUTP14A<sup>min</sup>) was inserted into MacroLab vector 1M (Addgene #29656) in fusion with an amino-terminal His<sub>6</sub>-MBP-tag. All DHX37 constructs, except MmDHX37<sup>CTD</sup> were expressed in Sf9 insect cells using the Bac-to-Bac system (Invitrogen) according to standard protocols. Briefly, Sf9 insect cells were infected with P3 virus (2.5 mL per 0.5 L) at a density of  $1.0 \times 10^6$  mL<sup>-1</sup>. Sixty-four hours after infection Sf9 cells were harvested and resuspended in Sf9 lysis buffer (20 mM Tris pH 8.0, 500 mM NaCl, 5 mM imidazole, 0.1% Tween 20, supplemented with Roche Protease Inhibitor Cocktail; 50 mL per 1 L of culture). Cells were lysed for 25 min by sonication (20% amplitude, 3 sec pulse on, 17 sec pulse off, 4°C) and lysates were clarified at 18,000g for 30 min at 4°C.

Full length His<sub>6</sub>-MmDHX37, His<sub>6</sub>-MmDHX37<sup>179–1150</sup>, and His<sub>6</sub>-MmDHX37<sup>179–1150</sup>/StreptII-GFP-HsUTP14A complex were purified by Nickel affinity chromatography (Ni<sup>2+</sup>-NTA-sepharose resin; Qiagen). The column was washed with Ni-wash buffer (20 mM Tris pH 8.0, 500 mM NaCl, 20 mM imidazole) and bound protein was eluted with Ni-elution buffer (20 mM Tris pH 8.0, 150 mM imidazole, 250 mM NaCl for MmDHX37<sup>179–1150</sup>; 20 mM Tris pH 8.0, 150 mM imidazole, 500 mM NaCl for MmDHX37 and the His<sub>6</sub>-MmDHX37<sup>179–1150</sup>/StreptII-GFP-HsUTP14A complex). The protein was subsequently cleaved with Tobacco Etch Virus (TEV) protease while dialyzed overnight at 4°C against 20 mM Tris pH 7.5, 250 mM NaCl (500 mM NaCl for MmDHX37 and complex),

1 mM TCEP pH 7.0 [Tris(2-carboxyethyl)phosphine hydrochloride]. In a second Ni-affinity chromatography step uncleaved His<sub>6</sub>-protein, His<sub>6</sub>-tagged TEV, and free His<sub>6</sub>-tag were removed. After the second Ni-affinity step, the NaCl concentration in the flow-through fraction was reduced to 250 mM using K0 buffer (20 mM Tris pH 7.5, 1 mM DTT) if necessary. The cleaved proteins were subjected to cation exchange chromatography using a Heparin column (HiTrap Heparin, GE Healthcare). The column was washed using IEX buffer A (20 mM Tris pH 7.5, 100 mM NaCl, 1 mM DTT) and protein was eluted by applying a gradient from 0% to 100% IEX buffer B (IEX buffer B: 20 mM Tris pH 7.5, 1000 mM NaCl, 1 mM DTT) over 20 column volumes. DHX37 was finally purified by a size exclusion chromatography step using a Superdex 200 column and eluted in 20 mM HEPES pH 7.5, 250 mM KCl (350 mM KCl for MmDHX37 and MmDHX37<sup>179–1150</sup>/HsUTP14A complex), 1 mM DTT. Eluted protein samples were concentrated and flash frozen with liquid nitrogen and stored at  $-80^{\circ}\text{C}$ .

GST-tagged MmDHX37<sup>CTD</sup> fusion protein was expressed in *E. coli* Rosetta 2 DE3 cells (Novagen, Merck Millipore) for 4 h at  $37^{\circ}\text{C}$  after induction with 0.2 mM IPTG (isopropyl- $\beta$ -D-thiogalactopyranoside). The protein was resuspended in lysis buffer (15 mL per 1 L culture; 20 mM Tris pH 8.0, 500 mM NaCl, 1 mM DTT) and lysed using a cell homogenizer (HPL6 cell homogenizer; passed three times,  $4^{\circ}\text{C}$ ). Cell lysates were clarified at 18,000g for 30 min at  $4^{\circ}\text{C}$ . GST-tagged MmDHX37<sup>CTD</sup> was purified by glutathione (GSH) affinity chromatography. The bound protein was washed with wash buffer (20 mM Tris pH 8.0, 500 mM NaCl, 1 mM DTT) and eluted in lysis buffer supplemented with 10 mM reduced GSH. MmDHX37<sup>CTD</sup> was finally purified by a size exclusion chromatography step using a Superdex 200 column and eluted in 20 mM HEPES pH 7.5, 250 mM KCl, 1 mM DTT. The protein was flash frozen with liquid nitrogen and stored at  $-80^{\circ}\text{C}$ .

His<sub>6</sub>-(StreptII)<sub>2</sub>-MmDHX37<sup>139–1150</sup> was purified by Nickel affinity chromatography as described above and dialyzed overnight at  $4^{\circ}\text{C}$  against 20 mM Tris pH 7.5, 250 mM NaCl, 1 mM DTT without tag cleavage. The protein was flash frozen with liquid nitrogen and stored at  $-80^{\circ}\text{C}$ . Catalytic inactive MmDHX37<sup>179–1150\_E369Q</sup> (MmDHX37<sup>EQ</sup>) construct was generated by site-directed mutagenesis (QuikChange) using 438B-MmDHX37<sup>179–1150</sup> as PCR template, and expressed and purified as His<sub>6</sub>-MmDHX37<sup>179–1150</sup>.

His<sub>6</sub>-MBP-HsUTP14A<sup>min</sup> fusion protein was expressed in Rosetta 2 DE3 cells (Novagen, Merck Millipore) for 4 h at  $37^{\circ}\text{C}$  after induction with 0.2 mM IPTG. The protein was resuspended in Ni-NTA lysis buffer (15 mL per 1 L culture; 20 mM Tris pH 8.0, 250 mM NaCl, 5 mM imidazole) and lysed using a cell homogenizer (HPL6 cell homogenizer; passed three times,  $4^{\circ}\text{C}$ ). Cell lysates were harvested and purified by Ni-NTA affinity chromatography as described above. Before flash freezing, the imidazole concentration of the protein sample was adjusted to 30 mM imidazole. The protein sample was stored at  $-80^{\circ}\text{C}$ . HsUTP14A alanine substitution mutant constructs, His<sub>6</sub>-MBP-HsUTP14A<sup>min\_MIDL</sup> and His<sub>6</sub>-MBP-HsUTP14A<sup>min\_MIEF</sup>, were generated using inverse PCR with 1M-HsUTP14A<sup>min</sup> as PCR template, and expressed and purified as His<sub>6</sub>-MBP-HsUTP14A<sup>min</sup>. To reconstitute MmDHX37<sup>179–1150</sup>/UTP14A<sup>min</sup> complex, a twofold molar excess of His<sub>6</sub>-MBP-HsUTP14A<sup>min</sup> was added to MmDHX37<sup>179–1150</sup> and incubated at room temperature for 10 min.

## Crystallization and data collection

Crystals of MmDHX37<sup>179–1150</sup> bound to U<sub>10</sub> RNA were grown by the hanging drop vapor diffusion method by mixing equal volumes (1  $\mu\text{L}$  + 1  $\mu\text{L}$ ) of MmDHX37<sup>179–1150</sup> (preincubated at a concentration of 7 mg mL<sup>-1</sup> with 1.5-fold molar excess of single-stranded U<sub>10</sub> RNA, 1 mM AMPPNP and 2.5 mM MgCl<sub>2</sub>), and reservoir solution containing 0.1 M Bis-Tris propane pH 7.0, 0.2 M NaNO<sub>3</sub>, 15% (w/v) polyethylene glycol (PEG) 3400 and incubation at  $20^{\circ}\text{C}$ . The size and morphology of crystals was optimized by microseeding. Au-derivatized crystals were obtained by soaking native crystals for 15 min in mother liquor supplemented with 10 mM K[Au(CN)<sub>2</sub>]. Before flash cooling in liquid nitrogen, crystals were cryoprotected by brief soaking in mother liquor supplemented with 15% (v/v) ethylene glycol. X-ray diffraction data were collected at beamline X06DA (PXIII) of the Swiss Light Source (Paul Scherrer Institute, Villigen, Switzerland). The crystals belonged to space group P2<sub>1</sub>, contained two copies of the DHX37-RNA complex in the asymmetric unit, and diffracted to a resolution of 2.9 Å (native crystal) or 3.0 Å (Au-derivatized crystal). For the Au-derivatized crystal a single-wavelength anomalous diffraction (SAD) experiment was performed. From a single crystal, seven 360° data sets were measured at a wavelength of 1.03980 Å, which corresponds to the Au L-III absorption edge. The diffraction data were processed using XDS (Kabsch 2010). SAD data was additionally scaled and merged with XSCALE (Kabsch 2010).

## Structure determination, refinement, and analysis

Structure determination was performed with the Phenix suite (Adams et al. 2010) based on the SAD data obtained from the Au-derivatized crystal. Twenty-seven Au sites were identified using phenix.hyss. Substructure refinement, phasing and density modification in phenix.autosol (Zwart et al. 2008) resulted in a clearly interpretable experimental electron density map, which was used for manual model building with COOT (Emsley and Cowtan 2004). The majority of DHX37 residues as well as uridine ribonucleotides could be assigned and built into the experimental electron density map. Next, molecular replacement was performed with the high-resolution native data set using phenix.phaser (McCoy et al. 2007). Structure refinement was carried out with phenix.refine (Afonine et al. 2012). For the initial refinement round, the B-factors of the starting model were set to 40 Å<sup>2</sup>. The initial round of refinement included rigid-body refinement, atomic coordinate refinement, and individual B-factor refinement. This was followed by iterative rounds of atomic coordinate, TLS and individual B-factor refinement, alternated with manual building in COOT to complete the atomic model. Noncrystallographic symmetry restraints were applied throughout the refinement and only released in the final round. The final atomic model was refined at a resolution of 2.9 Å with an  $R_{\text{free}}$  of 29.7% and an  $R_{\text{work}}$  of 26.8%. X-ray crystallographic data collection and refinement parameters are provided in Table 2. The final model of the DHX37-U<sub>10</sub> complex comprises 10 uridine ribonucleotides and DHX37 amino acid residues 231–1149, except for residues 432 and 490–578, which are disordered in electron density maps. Analysis of the two MmDHX37<sup>179–1150</sup> molecules using



**TABLE 2.** Data collection and refinement statistics

Data set	DHX37	
	Native	Hg SAD
X-ray source	SLS X06DA (PXIII)	SLS X06DA (PXIII)
Space group	$P2_1$	$P2_1$
Cell dimensions		
<i>a</i> , <i>b</i> , <i>c</i> (Å)	83.83, 137.60, 94.23	83.69, 138.16, 94.89
$\alpha$ , $\beta$ , $\gamma$ (°)	90.00, 93.34, 90.00	90.00, 93.67, 90.00
Wavelength (Å)	1.0077	1.0398
Resolution (Å) <sup>a</sup>	47.03–2.88 (2.98–2.88)	45.61–2.99 (3.10–2.99)
$R_{\text{merge}}$ <sup>a</sup>	0.112 (0.684)	0.186 (2.054)
$CC_{1/2}$ <sup>a</sup>	0.996 (0.746)	0.999 (0.761)
$I/\sigma I$ <sup>a</sup>	13.65 (2.17)	29.47 (3.04)
Observations <sup>a</sup>	320,411 (25,685)	2,381,443 (213,156)
Unique reflections <sup>a</sup>	47,756 (4209)	43,382 (4240)
Multiplicity <sup>a</sup>	6.7 (6.1)	54.9 (50.3)
Completeness (%) <sup>a</sup>	98.52 (87.65)	99.78 (98.38)
Refinement		
Resolution (Å)	47.03–2.88	
No. reflections	47,734 (4209)	
$R_{\text{work}}/R_{\text{free}}$	0.270/0.295	
No. atoms		
Protein	12,605	
RNA	394	
B-factors (Å <sup>2</sup> )		
Mean	65.3	
Protein	65.2	
RNA	68.2	
R.m.s. deviations		
Bond lengths (Å)	0.005	
Bond angles (°)	0.79	
Ramachandran plot		
% Favored	96.8	
% Allowed	3.2	
% Outliers	0.0	

<sup>a</sup>Values in parentheses denote highest resolution shell.

DynDom (Hayward et al. 1997) indicates that the major conformational difference between the two copies in the asymmetric unit involves a rotation by 8.8° of the CTD relative to the helicase core (comprising domains RecA1, RecA2, HA2, and OB) about a pivot point centered on residues 1007–1008. The helicase cores of the two DHX37<sup>179–1150</sup> copies superimpose as rigid bodies and align with an RMSD of 1.03 Å over 659 Cα atoms. The CTD domains superimpose with an RMSD of 0.45 Å over 127 Cα atoms. Structure superpositions of DHX37 and Prp43 and DHX36 were performed using the DALI server (Holm and Sander 1995). To model the conformation of MmDHX37 in the activated ATP- and RNA-bound state (MmDHX37-ATP-RNA complex), individual DHX37 domains (as defined in Fig. 1A), except for the CTD,

were individually superimposed with the structure of the crPrp43-RNA-ADP.BeF<sub>3</sub><sup>−</sup> complex (Tauchert et al. 2017) and combined in a single-polypeptide model. Domain motions predicted to occur during the RNA translocation cycle were subsequently analysed by comparing the resulting model with the crystal structure of the nucleotide-free MmDHX37-RNA complex in DynDom.

### ATPase activity assays

The ATPase activities of DHX37 constructs in the presence and absence of UTP14A were measured using the EnzCheck

Phosphate kit (Invitrogen) at 26°C. Here, the provided 2-amino-6-mercapto-7-methylpurine riboside (MESG) substrate together with the released inorganic phosphate ( $P_i$ ) is enzymatically converted to 2-amino-6-mercapto-7-methyl-purine which absorbs light at 360 nm, thus, facilitating the measurement of the  $P_i$  concentration in the reaction mixture. All reactions were performed in triplicates in a buffer containing 50 mM HEPES pH 7.5, 150 mM KCl, 1 mM DTT, and 1 mM  $MgCl_2$ , in a total volume of 200  $\mu$ L. For ATPase activity assays (Supplemental Figs. S1A, S9B,D, S11A) 0.25  $\mu$ M DHX37 (and complexes) was used. For reactions carried out in the presence of RNA, 0.25  $\mu$ M single-stranded  $U_{10}$  RNA was added to the protein samples, and the samples were incubated for 10 min at room temperature. Reactions were supplemented with 0.2 U purine nucleoside phosphorylase (PNP) and 1 mM MESG according to the manufacturer's protocol. The ATPase reactions were initiated by adding ATP and  $MgCl_2$  (both at 2 mM final concentration) to the samples. Absorbance at 360 nm was measured once per minute over 40 min in a microplate reader (PheraStar FSX fluorescence plate reader; BMG Labtech). The absorbance values at 360 nm were corrected for background absorbance. To quantify the released  $P_i$ , a calibration curve was generated with the phosphate standard provided together with the kit. The resulting values were plotted as a function of time using the GraphPad Prism software. To compare ATPase activities between samples, data points within the linear range were fitted with linear regression, and listed in Supplemental Table S1. The data represent mean values from three replicates, while the error bars indicate the standard error.

For Michaelis–Menten analysis (Fig. 3C), MmDHX37<sup>179–1150</sup> was used at final concentrations of 2  $\mu$ M in the absence of RNA or 1  $\mu$ M when assayed in the presence of  $U_{10}$  RNA. The MmDHX37<sup>179–1150</sup>/UTP14A<sup>min</sup> complex was reconstituted as described above and assayed at a final concentration of 0.125  $\mu$ M. For experiments carried out in the presence of RNA, a 1.5-fold molar excess of  $U_{10}$  RNA was added. ATP was used at concentrations of 7.8125–1000  $\mu$ M. ATPase activities were normalized for DHX37 concentrations to allow direct comparisons. The data represent mean values from three replicates, while the error bars indicate the standard error.  $K_M$  and  $k_{cat}$  values were fitted according to the Michaelis–Menten equation in GraphPad Prism and are provided in Table 1.

To investigate the effect of RNA on ATPase activity (Fig. 3D), MmDHX37<sup>179–1150</sup> and the MmDHX37<sup>179–1150</sup>/UTP14A<sup>min</sup> complex were used at a final concentration of 0.5  $\mu$ M, ATP at a final concentration of 1 mM, and  $U_{10}$  RNA was titrated to final concentrations of 0.1–12.8  $\mu$ M. Absorbance values at 360 nm were recorded every 15 sec for 25 min and quantified using the phosphate standard as described above. Initial reaction rates were calculated from the linear region of the absorbance curve. The resulting values were plotted as a function of ATP concentration (or RNA concentration) using GraphPad Prism and fitted by nonlinear regression analysis using the dose-versus-response relationship to calculate half-maximal effective concentration ( $EC_{50}$ ) values.

## Equilibrium RNA binding experiments

RNA binding affinities of DHX37 in the presence and absence of UTP14A were determined by FP measurements with 5'-

ATTO532-labeled  $U_{12}$  RNA at 26°C using a PheraStar FSX fluorescence plate reader. All reactions were carried out in triplicates in a buffer containing 20 mM HEPES pH 7.5, 150 mM KCl, 2 mM  $MgCl_2$ , and 0.1% Tween20, in a total volume of 50  $\mu$ L. NaCl,  $MgCl_2$ , and Tween20 concentrations of the protein samples were adjusted to 150 mM, 2 mM, and 0.1% Tween20, respectively, using 1.5x K50 buffer (20 mM HEPES pH 7.5, 50 mM KCl, 3 mM  $MgCl_2$ , and 0.15% Tween20. MmDHX37<sup>179–1150</sup>/UTP14A<sup>min</sup> complex was reconstituted as described above. For the RNA binding measurements, 0.25 nM Atto<sup>532</sup>-labeled  $U_{12}$  RNA were titrated with MmDHX37<sup>179–1150</sup> or MmDHX37<sup>179–1150</sup>/UTP14A<sup>min</sup> complex at varying concentrations ranging from 0.27 nM to 17.5  $\mu$ M. For measurements performed in the presence of adenosine nucleotides, ADP and AMPPNP were added at final concentrations of 1 mM. The AMPPNP stock (Jena Bioscience) was analysed by HPLC prior to its usage and estimated to be >95% pure. Fluorescence polarization (FP) signal was measured with an excitation filter with a central wavelength of 540 nm, and P and S emission filters with a central wavelength of 590 nm. The resulting FP values were plotted using the Graphpad Prism software as a function of concentration and fitted to a modified one-site binding equation accounting for ligand depletion:

$$Y = i + \left( (a - i) \cdot \frac{(c + K + X) - \sqrt{((c + K + X)^2 - 4cX)}}{2c} \right),$$

where  $K$  is the  $K_d$  to be determined,  $c$  is the total concentration of Atto532-labeled RNA,  $i$  is the  $Y_{min}$  value, and  $a$  is the  $Y_{max}$  value. The polarization amplitudes were normalized to 1. Error bars indicate standard errors of the mean (SEM) for five consecutive measurements of a single representative sample. The calculated  $K_d$  values are provided in Table 1.

## RNA pull-down experiment

For the RNA binding assay, 3  $\mu$ g of MmDHX37<sup>179–1150</sup> were incubated in binding buffer (20 mM HEPES pH 7.5, 150 mM KCl, 2 mM  $MgCl_2$ , 0.1% Tween20, 1 mM DTT) in the presence or absence of 1 mM adenosine nucleotides (AMPPNP or ADP) and 333 nM 5' end biotinylated  $U_{20}$  single-stranded RNA in a volume of 30  $\mu$ L. After 60 min incubation at 4°C, 50  $\mu$ L magnetic streptavidin coupled beads (Streptavidin-coupled Dynabeads, ThermoScientific) and 200  $\mu$ L of binding buffer (supplemented with or without 1 mM AMPPNP and ADP) were added. The samples were incubated with the beads for 1 h at 4°C. Beads were washed three times with 0.5 mL binding buffer and samples were eluted in SDS dye. Analysis of the eluted samples was done by SDS-PAGE.

## DHX37-UTP14A pull-down experiments

HEK293T cells were grown in DMEM supplemented with 10% fetal bovine serum, 100 U  $mL^{-1}$  penicillin/streptomycin (Sigma), 2 mM L-glutamine (Sigma) and 100  $\mu$ g  $mL^{-1}$  sodium pyruvate at 37°C and 5%  $CO_2$ . For overexpression in mammalian HEK293T cells, full-length HsUTP14A and truncations were PCR amplified and were cloned into a homemade plasmid based on the ligation-independent MacroLab vector 6A (Addgene #30124),

adding an amino-terminal His<sub>6</sub>-(HA)<sub>3</sub>-mCherry-tag. HsUTP14A<sup>min</sup> multi-alanine binding mutants M558A/I559A/D560A/L561A, (HsUTP14A<sup>min</sup><sub>MIDL</sub>) and M594A/I595A/E597A/F599A (HsUTP14A<sup>min</sup><sub>MIEF</sub>) constructs were generated by inverse PCR. Transfections were performed in 10-cm dishes seeded with  $2 \times 10^6$  cells using the X-tremeGENE HP DNA Transfection Reagent (Roche) and 3–4.5 µg plasmid DNA per transfection. HEK293T cells were collected 48 h post-transfection, washed twice with PBS and resuspended in 1.5 mL assay buffer (20 mM Tris pH 7.5, 150 mM KCl, 1 mM EDTA, 0.1% [v/v] Tween20, 1 mM DTT) supplemented with Roche Protease Inhibitor Cocktail. The cells were detached by gentle rocking at 4°C for 10 min and lysed by sonication (10% amplitude, 0.5 sec pulse on, 2 sec pulse off, 15 times, 4°C). The lysates were cleared by two centrifugation steps at 14,000 rpm for 15 min and incubated with 5 µg/mL RNase A (Thermo Fisher) for 20 min on ice. Per reaction, 25 µL (packed volume) Strep-Tactin Sepharose beads (IBA) were incubated with 250 µg of His<sub>6</sub>-(StrepII)-MmDHX37<sup>139–1150</sup> for 30 min at 4°C and washed three times with HEK assay buffer. For the pull-downs, 450 µL of cleared HEK293T cell lysates were incubated with 25 µL MmDHX37-Strep-beads for 1 h at 4°C. For the no bait control, 25 µL Strep-beads without MmDHX37 were incubated with 450 µL HEK293T cell lysate. The beads were washed four times with HEK293T assay buffer and bound proteins were eluted in two 10 min-steps on ice with 15 µL elution buffer (20 mM Tris pH 8.0, 150 mM KCl, 1 mM EDTA, 2.5 mM desthiobiotin) each. The eluted samples and the input lysate were loaded on SDS-PAGE without prior boiling to avoid mCherry denaturation, visualized on a Typhoon FLA9500 fluorescence scanner (GE Healthcare) and subsequently stained with Coomassie brilliant blue R250.

For MBP pull-down experiments, 10 µg of MBP-tagged UTP14A<sup>min</sup> were immobilized on 50 µL (50% slurry) of amylose resin (NEB) in a binding buffer containing 20 mM Tris pH 7.5, 150 mM KCl, 1 mM DTT and 0.1% Tween 20. The beads were washed and incubated with 15 µg of GST-tagged MmDHX37<sup>CTD</sup> in binding buffer for 1 h at 4°C with gentle rocking. Subsequently the beads were washed four times with 1 mL of binding buffer and proteins were eluted by boiling the resin with 20 µL of SDS-PAGE sample buffer. Proteins were analyzed by SDS-PAGE and visualized by Coomassie staining. For GST pull-down experiments, 10 µg of GST-tagged MmDHX37<sup>CTD</sup> were immobilized on 50 µL (50% slurry) of glutathione resin (GE Healthcare) in binding buffer. The beads were washed and incubated with 15 µg of MBP-tagged HsUTP14A<sup>555–619</sup> in binding buffer for 1 h at 4°C with gentle rocking. The beads were washed four times with 1 mL of binding buffer and proteins were eluted by boiling the resin with 20 µL of SDS-PAGE sample buffer. Proteins were analyzed by SDS-PAGE and visualized by Coomassie staining.

### DHX37 plasmid constructs for mammalian cell expression

For expression in mammalian cells (HEK293 and HeLa FlpIn TREx cells), full-length MmDHX37 and MmDHX37 encompassing residues 2–1003 (MmDHX37<sup>ACTD</sup>) were PCR-amplified and subcloned into a modified pcDNA5/FRT/TO expression vector, adding an amino-terminal EGFP-tag (EGFP-MmDHX37<sup>WT</sup> and EGFP-MmDHX37<sup>ACTD</sup>) or an amino-terminal HA-(Strep)<sub>2</sub>-tag

(HAsT-MmDHX37<sup>ACTD</sup>). The EGFP fragment was inserted into the KpnI/HindIII sites of pcDNA5<sup>TM</sup>/FRT/TO (Invitrogen). Catalytically inactive MmDHX37 was generated by PCR using EGFP-MmDHX37<sup>WT</sup> as PCR template (EGFP-MmDHX37<sup>EQ</sup>).

### DHX37<sup>ACTD</sup>-UTP14A pull-down experiments

Transfections of HEK293T cells were performed as described above. Briefly, 3 µg plasmid DNA (EGFP-MmDHX37<sup>WT</sup> and EGFP-MmDHX37<sup>ACTD</sup>) were transfected in HEK293T cells at a density of  $2 \times 10^6$  cells using 9 µL of the X-tremeGENE HP DNA Transfection Reagent (Roche). Forty-eight hours posttransfections, cells were harvested and lysed in 1.5 mL assay buffer (20 mM Tris pH 7.5, 150 mM KCl, 1 mM EDTA, 0.1% [v/v] Tween20, 1 mM DTT) supplemented with Roche Protease Inhibitor Cocktail, in the presence of RNase A (Thermo Fisher). Per reaction, 50 µL (packed volume) Amylose beads (NEB) were incubated with 2 mg of recombinant His<sub>6</sub>-MBP-HsUTP14A<sup>min</sup> for 30 min at 4°C and washed three times with HEK assay buffer. For the pull-downs, 500 µL of cleared HEK293T cell lysates were incubated with 50 µL His<sub>6</sub>-MBP-HsUTP14A<sup>min</sup>-Amylose-beads for 1 h at 4°C. For the no bait control, 50 µL Amylose-beads without His<sub>6</sub>-MBP-HsUTP14A<sup>555–619</sup> were incubated with 500 µL HEK293T cell lysate. The beads were washed four times with HEK293T assay buffer and bound proteins were eluted in 1× SDS-PAGE sample loading dye supplemented with 10 mM maltose. Proteins were separated by SDS-PAGE with no prior boiling to avoid EGFP denaturation, visualized on a Typhoon FLA9500 fluorescence scanner (GE Healthcare) and subsequently stained with Coomassie brilliant blue R250.

### Expression and purification of DHX37<sup>ACTD</sup>

HAsT-MmDHX37<sup>ACTD</sup> was expressed in suspension-adapted HEK293S-GnTI<sup>−</sup> cells (ATCC CRL-3022). Cells were grown in HyClone TransFx-H media (Sigma) supplemented with 2 mM L-glutamine (Sigma), 100 U mL<sup>−1</sup> penicillin/streptomycin (Sigma), 1% FBS (Sigma), and 1% Pluornic F-68 (Thermo Fisher) at 37°C and 5% CO<sub>2</sub> while shaking at 185 rpm in TubeSpin Bioreactor 600 vessels. One day prior to transfection, cells were diluted to a density of  $0.7 \times 10^6$  cells mL<sup>−1</sup>. Transfection mix was prepared by diluting plasmid DNA (1.3 µg DNA per 10<sup>6</sup> cells) to 0.015 µg µL<sup>−1</sup> in DMEM media, and mixing it with 25 kDa linear polyethylenimine (PEI, Polysciences) at a concentration of 0.38 µg µL<sup>−1</sup>. After incubation for 10 min, the DNA-PEI complexes were diluted 10-fold into the cultured cells, and valproic acid (Sigma) was added to a final concentration of 3 mM. Forty-five hours post-transfection, cells were harvested by centrifugation at 500g for 10 min and washed with PBS. The cells were resuspended in lysis buffer (20 mM Tris pH 7.5, 350 mM KCl, 1 mM DTT, 0.05% Tween20, supplemented with Roche Protease Inhibitor Cocktail; 100 mL per 1 L of culture). Cells were lysed for 30 min by sonication (20% amplitude, 3 sec pulse on, 17 sec pulse off, 4°C) and lysates were clarified at 18,000g for 30 min at 4°C. HAsT-MmDHX37<sup>ACTD</sup> was purified by Strep-Tactin affinity chromatography using 2 mL of Strep-Tactin resin (Strep-Tactin Superflow; IBA). The bound protein was washed with lysis buffer (20 mM Tris pH 7.5, 350 mM KCl, 1 mM DTT, 0.05% Tween20)

and eluted with lysis buffer supplemented with 2.5 mM desthiobiotin.

### DHX37 overexpression experiments

HeLa FlpIn TReX cells (kind gift from Dr. M. Beck, EMBL, Heidelberg) were grown in DMEM containing 10% fetal bovine serum supplemented with penicillin/streptomycin at 37°C and 5% CO<sub>2</sub>. Monoclonal HeLa cell lines expressing amino-terminally EGFP-tagged WT, ΔCTD and E369Q MmDHX37 proteins were generated by stable integration of the plasmid DNA constructs into the FRT site. All cell lines were tested negative for mycoplasma using PCR-based testing. Expression of the constructs was induced by the addition of tetracycline for 24 h (WT: 5.6 ng/mL, ΔCTD: 500 ng/mL, E369Q: 16.7 ng/mL).

### CRISPRi knockdown-rescue experiments

For CRISPRi-mediated DHX37 depletion, we modified the method originally described by the Weissman laboratory (Gilbert et al. 2014) by adding an additional nuclear localization signal to the catalytically inactive Cas9<sup>D10A, H840A</sup> mutant (dCas9). In brief, KRAB-dCas9 was amplified from pHR-SFFV-KRAB-dCas9-P2A-mCherry (kind gift of Dr. J. Weissman; Addgene, #60954). The resulting PCR product KRAB-dCas9-HA-2xSV40-NLS was cloned into the AgeI/XhoI sites of the pC2P vector (kind gift of Dr. M. Bühler; Friedrich Miescher Institute for Biomedical Research, Basel) containing a puromycin resistance cassette separated by a self-cleaving P2A (porcine teschovirus-1 2A) site from the upstream dCas9 fusion protein. Subsequently, the coding sequence for the importin-β binding (IBB) domain of importin α (RCH1) was inserted into the AgeI/HindIII-digested vector at the 5' end of the dCas9 cassette. A guide RNA targeting the DHX37 gene was inserted into the BsaI-sites of the same vector by primer annealing (5'-caacgGGTGGTCTGGGAACCCACGT-3' and 5'-aaacACGTGGGTTCCTCCAGACCACC), allowing sgRNA expression from a U6 promoter. An analogous vector containing a gRNA targeting the *Danio rerio tia1l* served as control (Lackner et al. 2015; 5'-caccgGGTATGTCTGGGAACCTCTCC-3' and 5'-aaacGGAGAGGTTCCCGACATACC). For CRISPRi rescue experiments, HeLa cell lines were transfected with CRISPRi vectors using the X-tremeGENE 9 transfection reagent (Roche). After 24 h, cells were reseeded and selected for successful transfection by treatment with 2 μg/mL puromycin for 24 h. Puromycin was washed out thoroughly with PBS and expression of the EGFP-MmDHX37 constructs was induced for 24 h. Ninety minutes before fixation, cells were treated with 20 nM leptomycin B (LMB, LC Laboratories) or solvent (ethanol).

### Immunoblot and immunofluorescence analysis and confocal microscopy

For protein analysis, whole cell extracts were prepared by lysis of cells in SDS sample buffer. Samples were analyzed by SDS-PAGE followed by wet-blotting onto PVDF membranes. Signals were detected on an Odyssey imaging system (LI-COR). The anti-ENP1 antibody has been described previously (Zemp et al. 2009). Commercial antibodies were anti-β-actin (A1987, Sigma)

and anti-DHX37 (ab70778, abcam). Immunofluorescence analysis was performed as described previously (Zemp et al. 2009). Images were acquired at a Zeiss LSM 880 upright microscope with a 63× 1.4 NA, oil, DIC Plan-Apochromat objective.

### DATA DEPOSITION

Coordinates and structure factors of the MmDHX37–RNA complex structure have been deposited in the Protein Data Bank with the entry code 6O16.

### SUPPLEMENTAL MATERIAL

Supplemental material is available for this article.

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